

1 IN THE UNITED STATES DISTRICT COURT  
 2 FOR THE NORTHERN DISTRICT OF OKLAHOMA

3 STATE OF OKLAHOMA, ex rel, )  
 4 W.A. DREW EDMONDSON, in his )  
 capacity as ATTORNEY GENERAL )  
 5 OF THE STATE OF OKLAHOMA, )  
 et al. )

6 Plaintiffs, )

7 V. ) No. 05-CV-329-GKF-SAJ  
 8 )

9 TYSON FOODS, INC., et al., )

10 Defendants. )

11  
 12  
 13 REPORTER'S TRANSCRIPT OF PROCEEDINGS

14 FEBRUARY 21, 2008

15 PRELIMINARY INJUNCTION HEARING

16 VOLUME III

17  
 18 BEFORE THE HONORABLE GREGORY K. FRIZZELL, Judge

19  
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**Exhibit 55**

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 UNITED STATES COURT REPORTER

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15 PROCEEDINGS

16 February 21, 2008

17 THE COURT: Mr. Bullock, Mr. George, and Ms.

18 Southerland and I spoke a second ago outside the courtroom with  
 19 regard to evidentiary matters. We've been going at such a  
 20 rapid pace and because there has been an agreement with regard  
 21 to exhibits on direct, there have been promises made to the  
 22 Court with respect to exhibits that have been used on cross  
 23 that they would be handled at the next break or at lunch that  
 24 has not been done. So the concern is that going forward, we  
 25 need to handle this matter very quickly or it presents real

1 A. Yes, we use the fecal indicator bacteria as a tracer or a  
2 surrogate to indicate the risk of the presence of human  
3 pathogens and thus the increased risk to human health from  
4 exposure to that water.

5 Q. Now, is it true that some pathogens that are in fecal  
6 material can be alive but not be culturable?

7 A. That's correct. The -- I guess the century old  
8 methodology for measuring bacterial concentrations is to  
9 culture them on some sort of an auger medium. We've known in  
10 the last 20 years or so that many organisms, when they're  
11 excreted from their host and they get out into the environment,  
12 may not die off, but they may become -- they may die off, but  
13 they may also become stressed, physiologically stressed, in  
14 which case they can no longer grow on the media that we  
15 normally use to culture them or detect them.

16 And so many studies have shown that when these  
17 bacteria become viable, we call this the viable but  
18 non-culturable phenomenon. They still have indications of  
19 metabolism and of the ability to sustain themselves. They can  
20 also be resuscitated or revived and start growing again when  
21 they get into a host, so when they get back into an environment  
22 that is conducive to their growth. So in spite of the fact  
23 that we cannot culture them and detect them, they still are  
24 potentially dangerous. And this is known in microbiology as  
25 the viable but non-culturable phenomenon. It's been seen in

1 so drying out. And again, it's very hard to say, it depends on  
2 a lot of common conditions that the bacteria encounter. If  
3 they are exposed fully to ultraviolet radiation and desiccated,  
4 it may take only a matter of hours for them to be permanently  
5 inactivated or killed. On the other hand, if they're shielded  
6 from radiation, if they're provided with some moisture, then  
7 they may persist for up to months at a time.

8 THE COURT: Thank you. Mr. Page.

9 MR. PAGE: Thank you, Your Honor.

10 Q. (By Mr. Page) So those bacteria can remain viable for  
11 months at a time if they have certain environmental conditions  
12 available?

13 A. That's correct.

14 Q. At the same time, if you use a standard method to try to  
15 identify that bacteria in the environment, it wouldn't  
16 necessarily be culturable?

17 A. That's correct, because the bacteria may be surviving and  
18 persisting in the environment, but they may be stressed to the  
19 point where they won't grow on this basically artificial growth  
20 substrate that we're providing them.

21 Q. Now, if a pathogen such as Campylobacter goes into this  
22 viable but not culturable state, can it then also remain as a  
23 hazard to human health?

24 A. Yes, studies have shown that viable but non-culturable  
25 organisms, when passed into a host such as perhaps if they were

1 Mr. Page.

2 MR. PAGE: Thank you, Your Honor.

3 Q. (By Mr. Page) Dr. Harwood, back to Exhibit 433. This is  
4 simply a summary of Oklahoma and U.S. EPA standards as they  
5 apply to recreational water quality uses; correct?

6 A. That is correct.

7 Q. That's bathing, swimming, splashing in the water; correct?

8 A. Right, correct.

9 Q. And I want to make sure this is clear. If someone is in  
10 water, bathing or swimming or splashing in the water, and the  
11 bacteria, any of those three bacteria, are at or above those  
12 levels, what does the EPA say about the expected sickness rate?

13 A. The EPA's guidelines and epidemiology studies and other  
14 epidemiology show that there is an increased risk of illness as  
15 levels above those standards rise. And the specific illness  
16 upon which most of these studies are based is gastroenteritis,  
17 so vomiting, diarrhea, nausea, cramps.

18 Q. How many people will get sick?

19 A. If the standards are right at that level, that's expected  
20 to be 8 individuals per thousand recreational water users and  
21 then it will go up from there. For example, if the E. coli  
22 concentrations increase about tenfold from this standard, then  
23 it's expected that the chance of getting ill will double.

24 Q. Thank you, Doctor. Now I'd like to turn your attention to  
25 State's Exhibit 434. Again, we have a blow-up on the tripod

1 consider high risk. And for example, poultry feces contain --  
2 are known to very frequently contain Salmonella and  
3 Campylobacter. These are so-called zoonotic pathogens which  
4 means that they're inhabitants of the animal gastrointestinal  
5 tract but they cause disease in humans. And in fact,  
6 Campylobacteriosis and Salmonellosis are among the most  
7 prevalent of both waterborne and foodborne diseases.

8 Q. Both Campylobacter and Salmonella, are they both present  
9 in poultry waste?

10 A. Yes, they are.

11 Q. What about E. coli, is that also a zoonotic bacteria?

12 A. Well, the pathogenic forms of E. coli are, such as E. coli  
13 0157:H7R, yes, zoonotic forms as well.

14 Q. I'd like now to draw your attention to State's Exhibit  
15 437. Dr. Harwood, could you identify this exhibit for the  
16 Court, please?

17 A. Yes, this exhibit is a graph that was prepared from data  
18 that was collected in the IRW from 2005 to 2007. And it shows  
19 the relationship between E. coli concentrations on the vertical  
20 axis and fecal coliform concentrations on the horizontal axis.  
21 And what this graph shows is that the relationship between fecal  
22 coliforms and E. coli in the vast majority of the IRW samples  
23 is nearly equivalent and very linear with a slope of about one.  
24 And so these are highly correlated. And with this sort of  
25 information then, we can feel comfortable about applying the



1 Q. Doctor, I want to now refer you to an exhibit that  
2 Dr. Teaf referred to a couple of days ago, State's Exhibit 406.  
3 Would you please remind us what information is shown on Exhibit  
4 406?

5 A. This is a map of the Illinois River Watershed. And these  
6 various color segments are those that have been designated  
7 impaired due to high indicator bacteria levels by the State of  
8 Oklahoma. At each of the dots are public access site points  
9 along tributaries in the Illinois River itself. And the red  
10 dots indicate sites where water quality standards were exceeded  
11 by indicator bacteria. So showing that, in fact, people who  
12 are using the water, they're putting in at these public access  
13 points for, as Dr. Caneday explained, for floating, swimming,  
14 canoeing, these people are being exposed to these elevated  
15 levels of indicator bacteria and thus at increased risk for  
16 illness.

17 Q. Now, does this information have any importance to you as a  
18 microbiologist with regard to evaluating the health risks  
19 associated with the Illinois River?

20 A. Yes, because we know that -- since we know -- so these  
21 aren't small ditches that nobody goes in, this is a scenic  
22 river. It is used -- it's an Oklahoma scenic river. It's  
23 widely used for recreation as was mentioned before. We know  
24 that literally thousands of people are being exposed to these  
25 high levels of bacteria and the increased health risk that's

1 represented by them.

2 Q. Thank you, Doctor. I want to switch gears on you a little  
3 bit again. Do you have an opinion with respect to the source  
4 of bacteria that has impaired the IRW?

5 A. Yes, I believe that a significant portion of that is  
6 contributed by land applied poultry litter.

7 Q. And do you have an opinion as to what would happen if  
8 poultry waste land application was stopped in the IRW?

9 A. Yes, I believe that over time the levels of bacteria would  
10 decline and that the human health risk would be decreased.

11 Q. Okay. Do you have any specific evidence, Doctor, that  
12 contribution of poultry litter to lands in the IRW has  
13 contaminated the waters of the IRW?

14 A. Yes, we used a reliable method called polymerase chain  
15 reaction or PCR to develop a poultry litter specific biomarker  
16 which we use as a tracer to follow the pathway of poultry -- of  
17 microbial contamination from poultry litter throughout the  
18 Illinois River Watershed.

19 Q. Would you just define briefly what a biomarker is?

20 A. A biomarker would be a biological component of some  
21 organism. In this case it's a bacterium and in this case the  
22 biological component is a gene fragment that we were able to  
23 detect by PCR and this bacterium is highly associated with  
24 chicken -- with contaminated chicken litter.

25 Q. Doctor, are there differences between the PCR method of

1 have the questions of fate and transport through the watershed.  
2 And we also have the question of there are things that we don't  
3 know about the relative rates of transport of pathogens  
4 compared to indicator bacteria and indicator bacteria and  
5 pathogens compared to the biomarker. So just because we don't  
6 detect it doesn't mean that there was never any poultry  
7 contamination there.

8 Q. Does the biomarker have a different life span in the  
9 environment than, for example, a chemical?

10 A. Well, a chemical might be expected to persist indefinitely  
11 until it gets used through biogeochemical cycling but because  
12 bacteria are biological organisms, they have a certain amount  
13 of persistence time in the environment, so they will not  
14 persist indefinitely over time.

15 Q. What type of samples were analyzed with the PCR method?

16 A. We analyze poultry litter samples. We analyze land  
17 applied soil samples or soil samples which received land  
18 application of poultry litter. We amplified edge of field  
19 samples which are basically direct runoff from fields that had  
20 received land application of poultry litter. Surface water  
21 samples, including Illinois River samples and tributary  
22 samples. And groundwater samples, including geoprobe samples  
23 and well samples, and also spring samples.

24 Q. From the samples you analyzed for litter, what were the  
25 results with the PCR marker?

1 A. All of the litter samples were positive for the biomarker,  
2 quantifiable with levels of biomarker over -- up to over a  
3 billion copies per gram.

4 Q. What about the land applied field samples, what were the  
5 biomarker results for that?

6 A. The land applied field samples were about 90 percent  
7 positive for the biomarker. And the maximum, around the  
8 maximum value for that was 10 million copies per gram.

9 Q. And what about edge of field, the next step in the path,  
10 what about those for biomarker?

11 A. Edge of field samples, about 50 percent positive and a  
12 maximum value of about 10 million per liter.

13 Q. And the same --

14 THE COURT: Doctor -- excuse me just a second, Mr.  
15 Page. You say you worked with Dr. Olsen with regard to  
16 sampling strategy and collection. To the uninitiated such as  
17 myself, the first question that jumps to mind as I tried to  
18 superimpose the location of the poultry houses to this map is  
19 that when we're talking about the area of recreational  
20 activity, there don't seem to be as many sampling stations, but  
21 rather that sampling is occurring in the area where these  
22 poultry houses are located and which raises fate and transport  
23 issues. I mean, to the extent that we are really focused here  
24 in this case about the public health concerns, it implicates  
25 fate and transport of these bacterium from the areas of highest

1 poultry house location.

2 Why is it that you and Dr. Olsen didn't select more?

3 I see that you have some green RNA results down here in the  
4 area just above Lake Tenkiller showing detectable, but not  
5 quantifiable. To the extent that we're focusing here to some  
6 extent on recreational activity and the public health  
7 repercussions or impact, why is it that you and Dr. Olsen  
8 didn't pick those locations as opposed to the locations closer  
9 to the poultry houses?

10 THE WITNESS: That would be -- when we were planning  
11 the sampling strategy, the focus was to find the pathway that  
12 would start basically at the poultry litter -- or find if there  
13 was a pathway that would start at the poultry litter houses and  
14 proceed --

15 THE COURT: From a scientific point of view.

16 THE WITNESS: Right.

17 THE COURT: I understand completely, sure.

18 THE WITNESS: Right. And then so, yeah, and I have to  
19 admit that, in fact, if I had looked at this map a couple of  
20 months ago, I wouldn't even have known where the important  
21 recreational water bodies were. It wasn't something that --  
22 demonstrating that hypothesis in particular wasn't the focus.

23 THE COURT: You're trying to make the link?

24 THE WITNESS: Yes, exactly.

25 THE COURT: Right, I understand. Go ahead, Mr. Page.

1 MR. PAGE: Thank you, Your Honor.

2 Q. (By Mr. Page) Did you detect the biomarker in surface  
3 water samples?

4 A. Yes, we did. We detected the biomarker in 43 and a half  
5 percent or so of surface samples at levels up to 100,000 per  
6 liter.

7 Q. What about groundwater samples?

8 A. We did detect it in some groundwater samples, two  
9 groundwater samples to be exact, and at a level up to 20,000  
10 per liter. And two out of 22 samples would be 9 percent.

11 Q. Now, a similar question to what the Judge just asked you.  
12 What does this information tell you, if anything, with regard  
13 to the distribution or pathway of poultry waste bacteria in the  
14 IRW?

15 A. Well, it demonstrates that the bacteria are following the  
16 pathway or that they have a transport pathway from the fields  
17 to the surface waters and also into the substratum into that  
18 karst, that fractured karst substratum which then allows them  
19 to appear in the groundwater and then be transported back  
20 upward into the spring systems.

21 Q. Let me draw your attention or if you would, to sample  
22 marked LAL5A on this exhibit. Can you identify that location  
23 for the Court, please?

24 A. Yeah, I think so. LAL5A is right about here. That's a  
25 soil sample and from a land applied field. That one had 4

1 Q. Does that mean the poultry waste biomarker co-varies with  
2 the indicator bacteria?

3 A. Correct.

4 Q. What is the chance of, let's say, a mistake in this  
5 analysis?

6 A. That would be, again, it's P less than .0001, so less than  
7 one in a thousand that this relationship occurred by chance.

8 Q. Now, Dr. Harwood, earlier I believe you stated an opinion  
9 concerning the importance of poultry waste as a contaminant, a  
10 bacterial contaminant in the IRW?

11 A. Correct.

12 Q. Would you please restate that opinion?

13 A. Yes, my opinion is that the poultry waste -- land  
14 application of poultry waste in the IRW is a major contributor  
15 to elevated indicator bacteria loads in the Illinois River  
16 Watershed in these waters.

17 Q. Now, what evidence did you use to reach this conclusion?

18 A. I used the weight of evidence approach which is what  
19 typically one does when investigating ecological questions. So  
20 rather than relying on one line of investigation, integrated  
21 numerous lines. So that would be starting out with -- and not  
22 in any particular order. But since we're talking about it, the  
23 widespread and quantifiable presence of the poultry litter  
24 biomarker and the evident pathway in terms of its concentration  
25 gradient from the litter to the fields to the edge of the field